Claim Am ndments:

Please amend claims 1, 2, 3, 11, 14-17 and 26, without prejudice or disclaimer, as follows:

- 1. (Currently amended) A method for decreasing interferences which result in inaccurate readings in serum or plasma containing assay samples of specific binding assays comprising adding an effective amount of a large, unconjugated polycation to serum or plasma containing assay samples during the specific binding assay for decreasing interferences in said assays.
- 2. (Currently amended) The method of claim 1 where<u>in</u> the large polycation has a molecular weight of 3,000 daltons or greater.
- 3. (Currently amended) The method of claim 1 where in wherein the large polycation is a polylysine, polyornithine, polybrene or MERQUAT dimethyldiallylammonium chloride.
- 4. (Original) The method of claim 3 wherein the large polycation comprises a polylysine with a molecular weight ranging between 5,200 and 11,200 daltons.
- 5. (Original) The method of claim 4 wherein the large polycation comprises polylysine with a molecular weight of 8,800 daltons.

- 6. (Original) The method of claim 1 wherein the specific binding assay measures thyroid stimulating hormone, free prostate specific antigen, alpha fetal protein, Hepatitis B core antibody, Hepatitis B surface antibody or human immunodeficiency virus.
- 7. (Original) The method of claim 1 wherein said specific binding assay is performed on a solid phase.
- 8. (Original) The method of claim 7 wherein said solid phase comprises paramagnetic microparticles.
- 9. (Currently amended) A method for decreasing interferences which result in inaccurate readings in serum or plasma containing assay samples of a thyroid stimulating hormone specific binding assay comprising adding a large, unconjugated polycation to serum or plasma containing assay samples during the thyroid stimulating hormone specific binding assay for decreasing interferences in said assays.
- 10. (Original) The method of claim 9 where the large polycation has a molecular weight of 3,000 daltons or greater.
- 11. (Currently amended) The method of claim 9 where in wherein the large polycation is a polylysine, polyornithine, polybrene or MERQUAT dimethyldiallylammonium chloride.

- 12. (Original) The method of claim 11 wherein the large polycation comprises a polylysine with a molecular weight ranging between 5,200 and 11,200 daltons.
- 13. (Original) The method of claim 12 wherein the large polycation comprises polylysine with a molecular weight of 8,800 daltons.
- 14. (Currently amended) A method for decreasing interferences which result in inaccurate readings in serum or plasma containing assay samples of a thyroid stimulating hormone specific binding assay comprising:
- a) forming a first complex by incubating a serum or plasma sample with paramagnetic microparticles coated with anti-β TSH antibody and an assay diluent which comprises a large, unconjugated polycation, for a time and under conditions which allow the thyroid stimulating hormone present in the sample to bind to the anti-β TSH antibody coated particles for decreasing interferences in said assays;
- b) forming a second complex by incubating the first complex with an acridinium labeled conjugate comprising an anti-α TSH antibody, for a time and under conditions which allow the conjugate to bind to the first complex;
 - c) creating a chemiluminescent reaction in the second complex; and

- d) measuring the chemiluminescent reaction as relative light units wherein the amount of thyroid stimulating hormone in the plasma or serum sample is directly related to the measured relative light units.
- 15. (Currently amended) A method for decreasing interferences which result in inaccurate readings in serum or plasma containing assay samples of a free prostate antigen specific binding assay comprising adding a large, unconjugated polycation to serum or plasma containing assay samples during the free prostate specific antigen specific binding assay for decreasing interferences in said assays.
- 16. (Currently amended) The method of claim 16 wherein the large polycation is a polylysine or polycrnitine polyornithine.
- 17. (Currently amended) A method for decreasing interferences which result in inaccurate readings in serum or plasma containing assay samples of a free prostate specific antigen specific binding assay comprising:
- a) forming a first complex by incubating a serum or plasma sample with paramagnetic microparticles coated with an antibody specific for free PSA and an assay diluent which comprises a large, unconjugated polycation, for a time and under conditions which allow the free PSA present in the sample to bind to the antibody coated particles for decreasing interferences in said assays;
- b) forming a second complex by incubating the first complex with an acridinium labeled conjugate comprising an anti-PSA antibody, for a time and under conditions which allow the conjugate to bind to the first complex;

- c) creating a chemiluminescent reaction in the second complex; and
- d) measuring the chemiluminescent reaction as relative light units wherein the amount of prostate specific antigen in the plasma or serum sample is directly related to the measured relative light units.

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- 26. (Currently amended) A method for decreasing interferences which result in inaccurate readings in serum or plasma containing assay samples of total prostate specific antigen specific binding assay comprising:
- a) forming a first complex by incubating a serum or plasma sample with paramagnetic microparticles coated with an antibody which binds both free and complexed PSA and an assay diluent which comprises a large, unconjugated polycation, for a time and under conditions which allow the PSA present in the sample to bind to the antibody coated particles for decreasing interferences in said assays;
- b) forming a second complex by incubating the first complex with an acridinium labeled conjugate comprising an anti-PSA antibody, for a time and under conditions which allow the conjugate to bind to the first complex;
 - c) creating a chemiluminescent reaction in the second complex; and
- d) measuring the chemiluminescent reaction as relative light units wherein the amount of prostate specific antigen in the plasma or serum sample is directly related to the measured relative light units.